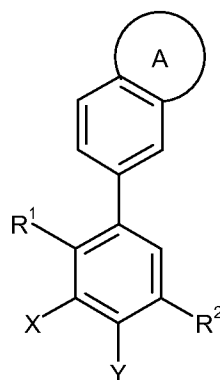


**Amendments to the claims**

This listing of claims will replace all prior versions, and listings, of claims in the application:

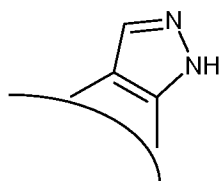
1. (Currently amended) A compound of formula (I):



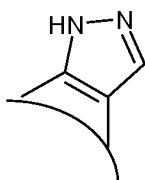
(I)

wherein

A is a fused 5-membered heteroaryl ring selected from



or



, which ring is

substituted by  $-(CH_2)_m$ heterocyclyl wherein the heterocyclyl is a 5- or 6-membered heterocyclic ring ~~containing one or two heteroatoms independently selected from tetrahydrofuranyl, tetrahydropyranyl or morpholinyl-oxygen, sulfur, and nitrogen~~ optionally substituted by up to two substituents independently selected from oxo,  $C_{1-6}$ alkyl,  $-(CH_2)_n$ phenyl,  $-OR^3$ ,  $-(CH_2)_nCO_2R^3$ ,  $-NR^3R^4$  and  $-CONR^3R^4$ , and

A is optionally further substituted by one substituent selected from  $-OR^3$ , halogen, trifluoromethyl,  $-CN$ ,  $-CO_2R^3$  and  $C_{1-6}$ alkyl optionally substituted by hydroxy;

$R^1$  is selected from methyl and chloro;

$R^2$  is selected from  $-NH-CO-R^5$  and  $-CO-NH-(CH_2)_q-R^6$ ;

$R^3$  and  $R^4$  are each independently selected from hydrogen and  $C_{1-6}$ alkyl;

$R^5$  is selected from hydrogen,  $C_{1-6}$ alkyl,  $-(CH_2)_q-C_{3-7}$ cycloalkyl, trifluoromethyl,  $-(CH_2)_r$ heteroaryl optionally substituted by  $R^7$  and/or  $R^8$ , and  $-(CH_2)_r$ phenyl optionally substituted by  $R^7$  and/or  $R^8$ ;

R<sup>6</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl, C<sub>3-7</sub>cycloalkyl, -CONHR<sup>9</sup>, phenyl optionally substituted by R<sup>7</sup> and/or R<sup>8</sup>, and heteroaryl optionally substituted by R<sup>7</sup> and/or R<sup>8</sup>;

R<sup>7</sup> is selected from C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, -(CH<sub>2</sub>)<sub>q</sub>-C<sub>3-7</sub>cycloalkyl, -CONR<sup>9</sup>R<sup>10</sup>, -NHCOR<sup>10</sup>, halogen, -CN, -(CH<sub>2</sub>)<sub>s</sub>NR<sup>11</sup>R<sup>12</sup>, trifluoromethyl, phenyl optionally substituted by one or more R<sup>8</sup> groups, and heteroaryl optionally substituted by one or more R<sup>8</sup> groups;

R<sup>8</sup> is selected from C<sub>1-6</sub>alkyl, C<sub>1-6</sub>alkoxy, halogen, trifluoromethyl, and -(CH<sub>2</sub>)<sub>s</sub>NR<sup>11</sup>R<sup>12</sup>;

R<sup>9</sup> and R<sup>10</sup> are each independently selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>9</sup> and R<sup>10</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>13</sup>, wherein the ring may be substituted by up to two C<sub>1-6</sub>alkyl groups;

R<sup>11</sup> is selected from hydrogen, C<sub>1-6</sub>alkyl and -(CH<sub>2</sub>)<sub>q</sub>-C<sub>3-7</sub>cycloalkyl optionally substituted by C<sub>1-6</sub>alkyl,

R<sup>12</sup> is selected from hydrogen and C<sub>1-6</sub>alkyl, or

R<sup>11</sup> and R<sup>12</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R<sup>13</sup>;

R<sup>13</sup> is selected from hydrogen and methyl;

X and Y are each independently selected from hydrogen, methyl and halogen;

m and q are each independently selected from 0, 1 and 2;

n and r are each independently selected from 0 and 1; and

s is selected from 0, 1, 2 and 3;

with the proviso that:

A is not substituted by -(CH<sub>2</sub>)<sub>m</sub>NR<sup>14</sup>R<sup>15</sup> wherein R<sup>14</sup> and R<sup>15</sup>, together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulphur and NR<sup>16</sup> wherein R<sup>16</sup> is hydrogen or methyl,

when m is 0, the -(CH<sub>2</sub>)<sub>m</sub>heterocyclyl group is not a 5- or 6-membered heterocyclyl ring containing nitrogen optionally substituted by C<sub>1-2</sub>alkyl or -(CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>3</sup>, and

the compound of formula (I) is not 1,1-dimethylethyl 4-(6-{5-[(cyclopropylamino)carbonyl]-2-methylphenyl}-1,2-benzisoxazol-3-yl)-1-piperazinecarboxylate;

or a pharmaceutically acceptable [[derivative]] salt thereof.

2. (Currently amended) A compound according to claim 1 wherein A is ~~a fused 5-membered heteroaryl ring containing two heteroatoms independently selected from oxygen and nitrogen~~ substituted by the -(CH<sub>2</sub>)<sub>m</sub>heterocyclyl moiety on the ring nitrogen of the A ring.

3. (Currently amended) A compound according to claim 1 wherein A is substituted by - (CH<sub>2</sub>)<sub>m</sub>heterocyclyl and [[wherein]] the heterocyclyl is an optionally substituted tetrahydropyranyl ~~a 5- or 6-membered ring containing one or two heteroatoms independently selected from oxygen and nitrogen optionally substituted by up to two substituents independently selected from oxo, C<sub>1-6</sub>alkyl, (CH<sub>2</sub>)<sub>n</sub>phenyl, OR<sup>3</sup>, (CH<sub>2</sub>)<sub>n</sub>CO<sub>2</sub>R<sup>3</sup>, NR<sup>3</sup>R<sup>4</sup> and CONR<sup>3</sup>R<sup>4</sup>.~~

4. (Previously Presented) A compound according to claim 1 wherein R<sup>1</sup> is methyl.

5. (Previously Presented) A compound according to claim 1 wherein R<sup>2</sup> is -CO-NH-(CH<sub>2</sub>)<sub>q</sub>-R<sup>6</sup>.

6. (Previously Presented) A compound according to claim 1 wherein X is fluorine.

7. (Currently amended) A compound according to claim 1 ~~substantially as hereinbefore defined with reference to any one of Examples 1 to 9, which is~~  
*N*-Cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2*H*-pyran-2-ylmethyl)-1*H*-indazol-5-yl]benzamide ;;  
*N*-Cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2-furanylmethyl)-1*H*-indazol-5-yl]benzamide ;  
3-{1-[(4-Benzylmorpholin-2-yl)methyl]-1*H*-indazol-5-yl}-*N*-cyclopropyl-5-fluoro-4-methylbenzamide ;  
*N*-Ethyl-4-methyl-3-[3-(tetrahydro-3-furanyl)-1*H*-indazol-6-yl]benzamide ;  
*N*-Ethyl-3-fluoro-4-methyl-5-[3-(tetrahydro-3-furanyl)-1*H*-indazol-6-yl]benzamide ;  
or a pharmaceutically acceptable [[derivative]] salt thereof.

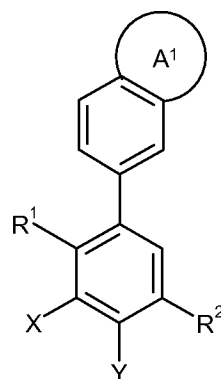
8. (Currently amended) A compound selected from:  
*N*-cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2*H*-pyran-2-ylmethyl)-1*H*-indazol-5-yl]benzamide;  
*N*-cyclopropyl-3-fluoro-4-methyl-5-[1-(tetrahydro-2-furanylmethyl)-1*H*-indazol-5-yl]benzamide; and  
3-{1-[(4-benzylmorpholin-2-yl)methyl]-1*H*-indazol-5-yl}-*N*-cyclopropyl-5-fluoro-4-methylbenzamide,  
or a pharmaceutically acceptable [[derivative]] salt thereof.

9. (Previously Presented) A pharmaceutical composition comprising at least one compound as claimed in claim 1, or a pharmaceutically acceptable derivative thereof, in association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.

10 -13 (Cancelled)

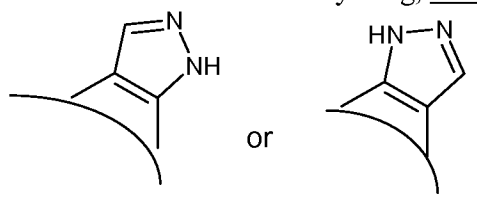
14. (Currently amended/Withdrawn) A process for preparing a compound of formula (I) ~~as claimed in~~ according to claim 1, or a pharmaceutically acceptable ~~[[derivative]]~~ salt thereof, which comprises:

(a) reacting a compound of formula (II)

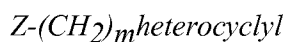


(II)

in which R<sup>1</sup>, R<sup>2</sup>, X and Y are as defined in claim 1 and A<sup>1</sup> is ~~an unsubstituted fused 5-membered heteroaryl ring~~ A is a fused 5-membered heteroaryl ring selected from



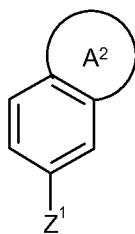
with a halide derivative of formula (III)



(III)

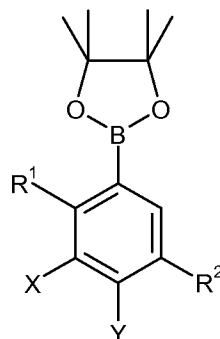
in which  $-(CH_2)_m \text{heterocyclyl}$  is as defined in claim 1 and Z is halogen, in the presence of a base;

(b) reacting a compound of formula (IV)

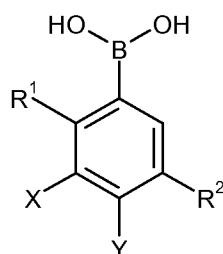


(IV)

in which  $A^2$  is A as defined in claim 1 or a protected form of A or  $A^1$ , and  $Z^1$  is halogen,  
with a compound of formula (VA) or (VB)



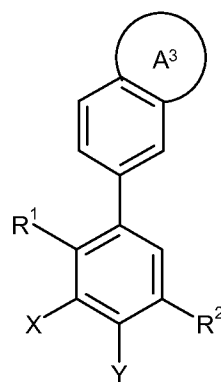
(VA)



(VB)

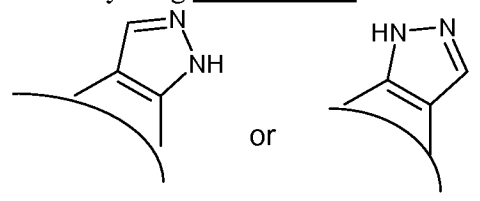
in which  $R^1$ ,  $R^2$ , X and Y are as defined in claim 1,  
in the presence of a catalyst;

(c) reacting a compound of formula (XI)



(XI)

in which  $R^1$ ,  $R^2$ ,  $X$  and  $Y$  are as defined in claim 1 and  $A^3$  is a fused 5-membered heteroaryl ring selected from



substituted by  $-(CH_2)_m$ heterocyclyl wherein the heterocyclyl is unsubstituted, with a suitable reagent; or

(d) final stage modification of one compound of formula (I) as defined in claim 1 to give another compound of formula (I) as defined in claim 1.

15. (Currently amended) A compound according to claim [[2]] 1 wherein A is substituted by  $-(CH_2)_m$ heterocyclyl and [[wherein]] the heterocyclyl is an optionally substituted tetrahydrofuranyl ~~a 5- or 6-membered ring containing one or two heteroatoms independently selected from oxygen and nitrogen optionally substituted by up to two substituents independently selected from oxo,  $C_{1-6}$ alkyl,  $(CH_2)_n$ phenyl,  $OR^3$ ,  $(CH_2)_nCO_2R^3$ ,  $NR^3R^4$  and  $CONR^3R^4$ .~~

16. (Previously Presented) A compound according to claim 15 wherein  $R^1$  is methyl.

17. (Previously Presented) A compound according to claim 15 wherein  $R^2$  is  $-CO-NH-(CH_2)_q-R^6$ .

18. (Previously Presented) A compound according to claim 15 wherein  $X$  is fluorine.

19. (new) A compound according to claim 1 wherein A is substituted by  $-(CH_2)_m$ heterocyclyl and the heterocyclyl is an optionally substituted morpholinyl.
20. (new) A compound according to claim 3 wherein m is 1.
21. (new) A compound according to claim 15 wherein m is 0.
22. (new) A compound according to claim 15 wherein m is 1.
23. (new) A compound according to claim 19 wherein m is 1.
24. (new) A compound according to claim 1 wherein A is substituted by the  $-(CH_2)_m$ heterocyclyl moiety on the 3-position of the A ring.
25. (new) A pharmaceutical composition comprising a compound according to claim 8, or a pharmaceutically acceptable salt thereof, in association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.